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         Feb 01
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                 frequency
NEWS
      5
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                 Gene Names now available in BIOSIS
NEWS
      6
NEWS
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                 TOXLIT no longer available
NEWS
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      9
         Mar 28
                 US Provisional Priorities searched with P in CA/CAplus
                 and USPATFULL
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         Mar 28
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NEWS 17
         Apr 22
                 BIOSIS Gene Names now available in TOXCENTER
NEWS 18
         Apr 22
                 Federal Research in Progress (FEDRIP) now available
NEWS EXPRESS
              February 1 CURRENT WINDOWS VERSION IS V6.0d,
              CURRENT MACINTOSH VERSION IS V6.0a(ENG) AND V6.0Ja(JP),
              AND CURRENT DISCOVER FILE IS DATED 05 FEBRUARY 2002
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=> s dentrit? and etch? 140 DENTRIT? 157940 ETCH? L1

6 DENTRIT? AND ETCH?

=> d all 1-6

ANSWER 1 OF 6 CA COPYRIGHT 2002 ACS

AN

ΤI Appearance form of iron compounds in ceramics during scanning electron microscope studies

ΑU Berger, Iris; Lange, Peter

Sekt. Baustoffverfahrenstech., Hochsch. Archit. Bauwes., Weimar, E. Ger. CS

SO Silikattechnik (1977), 28(11), 332-4 CODEN: SITKA7; ISSN: 0037-5233

DTJournal

German LA

CC 57-7 (Ceramics)

AB SEM (scanning electron microscopy) examns. of ceramics with added Fe and Fe compds. obtained under various heating and oxidative conditions were used to identify the type of reaction, if any, between the ceramic and the Fe and the changes occurring in the Fe and ceramics; the Fe remained unchanged at 1100.degree.. At 1200-1300.degree., the Fe somewhat penetrates into the surrounding ceramic, but without reaction. At 1400.degree., magnetite [1309-38-2] is formed, with a dentritic magnetite zone of peneration. At higher temps., magnetite dentritic formation is readily identified, even without prior etching of sample. Magnetite is formed under reducing conditions as isometric zonal crystals even at <1200.degree.. Other reaction products of Fe are also obsd. by SEM.

ST iron reaction ceramic magnetite

ΙT Ceramic materials and wares

(reaction of, with iron in situ)

IT 1309-38-2P, preparation

> RL: PEP (Physical, engineering or chemical process); PREP (Preparation); PROC (Process)

(formation of, in ceramics in reactions with iron)

IT 7439-89-6P, preparation

```
RL: PEP (Physical, engineering or chemical process); PREP (Preparation);
     PROC (Process)
        (formation of, in iron reaction in ceramics)
     ANSWER 2 OF 6 CA COPYRIGHT 2002 ACS
L1
AN
     78:114443 CA
     Crystallization of metal powders obtained by liquid-phase pulverization
TI
     Petrov, A. K.; Miroshnichenko, I. S.; Parabin, V. V.; Parabina, G. I.;
ΑU
     Sergeev, G. N.; Orlov, Yu. G.; Golovko, V. A.; Brekharya, G. P.
     Ukr. Nauchno-Issled. Inst. Spets, Stalei, Splavov Ferrosplavov, USSR
CS
SO
     Porosh. Met. (1973), 13(1), 16-20
     CODEN: PMANAI
DT
     Journal
LΑ
    Russian
     56-3 (Nonferrous Metals and Alloys)
CC
     Crystn. of pulverized molten metals was studied with emphasis on the
AB
     relation between the cooling rate and the formation of dendritic
     structures and the effect of crystn. parameters on the oxidn. of
     solidifying particles. Initial materials were molten R18 (Cr-W-V steel),
     Kh18N15 (Cr-Ni steel), El437B (ni-Cr alloy), and VK4A. Metallog,
     specimens were prepd. by compacting pulverized metal in brass casings, by
     mixing pulverized particles with a solidifying resin, and by pressing the
     power with pwd. Ni (1:4 ration). The last procedure made it possible to
     measure the oxide-layer thickness and to etch the specimens
     electrolytically. The structure of all the powders examd. is
     dentritic and strongly affected by the cooling rate, Vc. The
     distance between secondary dentritic branches can be used for
     characterizing Vc. This relation was confirmed by expts. with different
     metals that solidified over a broad cooling-rate range (10-107
     degrees/sec). When the dendritic parameter is plotted vs. Vc on a
     logarithmic scale, a linear dependence was found. This parameter is also
     dependent on the steel compn., but to a lesser extent. The mean thickness
     of the oxide layer on the surface of R18 particles decreases with Vc.
     Consequently, the oxidn. of pulverized particles takes place mainly during
     their crystn.; addnl. oxidn. of solid particles is negligible.
ST
     metal powder crystn; steel alloy powder crystn; nickel alloy powder
     crystn; chromium alloy powder crystn; tungsten steel powder crystn;
     vanadium steel powder crystn
IT
     Crystallization
        (of metal powders during atomization)
IT
     Atomization
        (of metal powders, crystn. in)
TT
                 37188-51-5 37241-62-6
                                          39391-97-4, uses and miscellaneous
     12719-18-5
     RL: USES (Uses)
        (atomization of, crystn. in)
    ANSWER 3 OF 6 CA COPYRIGHT 2002 ACS
L1
     69:81748 CA
ΑN
TI
    Method of etching silicon carbide
IN
    Haga, Leigh J.; Tucker, Thomas N.
PA
    Dow Corning Corp.
SO
     U.S., 2 pp.
     CODEN: USXXAM
DT
    Patent
LA
    English
NCL
    156017000
     71 (Electric Phenomena)
CC
FAN.CNT 1
     PATENT NO.
                     KIND DATE
                                          APPLICATION NO. DATE
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                                          -----
PΙ
    US 3398033
                  Α
                           19680820
                                          US 1965-435674
                                                           19650226
    A method of removing SiC from a Si surface by etching is
     described. The SiC is heated and then exposed to a halogen-O mixt.
     mixt. causes deterioration of the SiC and tends to make it porous.
```

Treatment by HF-HNO3 etching removes the remaining SiC from the Si after the SiC has become porous. E.g., a section of dentritic web Si crystal with a 1 mil SiC coating is placed in a reaction chamber and heated to 1200.degree.. An etch mixt. of Cl 25 and O 75 vol. % is fed into the chamber at 500 cc./min. for 2 hrs. The SiC layer at this point is deteriorated and porous. The surface is then treated with a mixt. of 1 part HF and 1 part HNO3 to remove the remaining SiC. etching Si carbides; carbides Si etching; silicon carbides etching Etching (of silicon carbide with halogen-oxygen mixts.) 7782-44-7, reactions RL: RCT (Reactant) (etching of silicon carbide (SiC) with mixts. of chlorine and fluorine and) 7782-41-4, reactions 7782-50-5, reactions RL: RCT (Reactant) (etching of silicon carbide (SiC) with mixts. of oxygen and) 409-21-2, reactions RL: RCT (Reactant) (etching of, with halogen-oxygen mixts.) ANSWER 4 OF 6 CA COPYRIGHT 2002 ACS 63:50719 CA OREF 63:9210h,9211a Transverse Hall coefficient and magnetoresistance of two-phase InSb-In Wieder, H. H.; Davis, N. M. U.S. Navy Ordnance Lab., Corona, CA Solid-State Electron. (1965), 8(7), 605-10 Journal English 9 (Electric and Magnetic Phenomena) The composite films recrystd. from the liquid phase are investigated as a function of the magnetic field before and after etching. The significance of the microgeometrical factors and the metallic inclusions is shown. The electron mobility in the InSb dentrites is estd. to be 3 .times. 104 cm.2/v.-sec. at 296.degree.K. Good agreement between theory and expt. is obtained. ANSWER 5 OF 6 CA COPYRIGHT 2002 ACS 55:1887 CA OREF 55:306i,307a-c Dentritic segregation in medium-alloyed structural steels Met., Sbornik Statei (1958), (No. 1), 115-34 Journal Unavailable 9 (Metallurgy) An app. was built for spectrographic microdetn. of Ni, Mn, Mo, etc., in segregates in steel ingots. Spark discharge (8-10 kv., 1000 cm.-0.01 .mu.f.) between 15-50-.mu. Pt or Cu wire and selected areas of the structure was used for excitation. The distance between the tip of the electrode and the specimen was 30 .mu.. The area of the specimen covered in the single detn. was .apprx.0.002 sq. mm. Cr 2663 and 2677, Ni 2416, and Mn 2933 A. lines were used. To facilitate the selection under a microscope of suitable portions of the specimen, dentritic structure was revealed by conventional etching, preceded by a heat-treatment in which transformation of austenite was stopped at the moment when it takes place only in the axes of dendrites. The detns. were carried out on open-hearth steels (duplex basic/acid) contq. Ni 2.5, Cr 1.9, and Mo 0.35%, and on standard- and vacuum-cast ingots for heavy rotors. With exptl. error of 4-6%, the av. contents of alloying elements

in interstitial segregates and in axes of dendrites were found to be the

ST

ΙT

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TT

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L1 AN

ΑU

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SO

DT

LA

CC

AΒ

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AN

ΑU SO

DT

LΑ

CC

AB

same in all parts of an ingot, except at the pipe. Differences of compn. in various parts of an ingot are only those between interstitial segregates and axes of dendrites. Segregation of C was evaluated indirectly by microhardness measurements and similar results were obtained.

ANSWER 6 OF 6 CA COPYRIGHT 2002 ACS L1 50:933 CA AN OREF 50:145c-d The application of radiographic methods to the structure studies of ΤI magnesium alloys Drits, M. E.; Sviderskaya, Z. A.; Kadaner, E. S. ΑU SO Zavodskaya Lab. (1955), 21, 831-3 Journal DTUnavailable LA CC 9 (Metallurgy and Metallography) AB The structure of Mg alloys was studied after the addn. of small amts. of a Ca isotope to the alloy, because the dentritic structure of Mg could not be revealed by etching. => s calix? and etch? 4628 CALIX? 157940 ETCH? L2 30 CALIX? AND ETCH? => s 12 not 11 30 L2 NOT L1 => d all 1=30 UNITS CONVERSION IS NOT AVAILABLE IN THE CURRENT FILE => d all 1-30 L3 ANSWER 1 OF 30 CA COPYRIGHT 2002 ACS AN 136:191684 CA Negative photoresist composition containing dendrimer and formation of ΤI photoresist pattern using it Afzali, Ardakani Ali; Breen, Tricia L.; Gelorme, Jeffrey Donald; Mitzi, IN David Brian; Lux, Michael Joseph PA International Business Machines Corp., USA Jpn. Kokai Tokkyo Koho, 12 pp. SO CODEN: JKXXAF DT Patent LA Japanese IC ICM G03F007-038 ICS H01L021-027 ·74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other

FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE ______ ---------PΙ JP 2002049152 A2 20020215 JP 2001-164184 20010531 PRAI US 2000-208269P P 20000531 US 2000-685414 Α 20001010 os MARPAT 136:191684 GΙ

Reprographic Processes)

Ι

The photoresist compn. comprises (1) aliph. dendrimer having .gtoreq.8 OH groups per mol., (2) glycoluril deriv. as crosslinker, (3) photoacid generator, and (4) org. solvent. Preferably, the dendrimer is calix [4]arene dendrimer deriv., the glycoluril deriv. has a general formula I (R1-2 = H, C1-6 alkyl, alkenyl, alkoxy, aryl; R3 = alkoxy), and the photoacid generator is onium salt. Photoresist pattern is formed by applying the compn. on a substrate, exposing to E-beam energy, baking, and developing with org. solvent. The resulting photoresist pattern is useful for etching semiconductors, ceramics, org. compds., and metals. The compn. has high sensitivity and resoln

ST neg photoresist dendrimer glycoluril deriv onium salt; sensitivity resoln neg photoresist calix arene dendrimer

IT Ceramics

Negative photoresists

Semiconductor materials

(solvent-developable neg. photoresist compn. contg. dendrimer, glycoluril deriv., and photoacid generator for high sensitivity and resoln.)

IT Dendritic polymers

RL: IMF (Industrial manufacture); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(solvent-developable neg. photoresist compn. contg. dendrimer, glycoluril deriv., and photoacid generator for high sensitivity and resoln.)

IT Metals, processes

RL: PEP (Physical, engineering or chemical process); PYP (Physical process); PROC (Process)

(solvent-developable neg. photoresist compn. contg. dendrimer, glycoluril deriv., and photoacid generator for high sensitivity and resoln.)

IT 57840-38-7, Triphenylsulfonium hexafluoroantimonate

RL: TEM (Technical or engineered material use); USES (Uses)
(UV 16974, photoacid generator; solvent-developable neg. photoresist compn. contg. dendrimer, glycoluril deriv., and photoacid generator for high sensitivity and resoln.)

IT 71443-52-2P

RL: IMF (Industrial manufacture); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(solvent-developable neg. photoresist compn. contg. dendrimer, glycoluril deriv., and photoacid generator for high sensitivity and resoln.)

IT 16837-14-2P

RL: PNU (Preparation, unclassified); RCT (Reactant); PREP (Preparation);
RACT (Reactant or reagent)

```
(solvent-developable neg. photoresist compn. contg. dendrimer,
        glycoluril deriv., and photoacid generator for high sensitivity and
        resoln.)
     50-00-0, Formaldehyde, reactions
                                        57-13-6, Urea, reactions
                                                                    3848-24-6,
IT
                     4767-03-7
                                  123830-85-3, Dimethoxypropane
     2,3-Hexanedione
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (solvent-developable neg. photoresist compn. contg. dendrimer,
        glycoluril deriv., and photoacid generator for high sensitivity and
        resoln.)
     220140-29-4
TT
     RL: TEM (Technical or engineered material use); USES (Uses)
        (solvent-developable neg. photoresist compn. contg. dendrimer,
        glycoluril deriv., and photoacid generator for high sensitivity and
        resoln.)
     ANSWER 2 OF 30 CA COPYRIGHT 2002 ACS
L3
AN
     135:303868 CA
     Syntheses and metal-ion binding properties of calix[4] arene
TT
     derivatives containing soft donor atoms: highly selective extraction
     reagents for Ag+
     Xie, Jian; Zheng, Qi-Yu; Zheng, Yan-Song; Chen, Chuan-Feng; Huang,
ΑU
     Zhi-Tang
     LMRSS, Center for Molecular Science, Institute of Chemistry, The Chinese
CS
     Academy of Sciences, Beijing, 100080, Peop. Rep. China
     Journal of Inclusion Phenomena and Macrocyclic Chemistry (2001), 40(1-2),
SO
     125-130
     CODEN: JIPCF5
     Kluwer Academic Publishers
PB
DT
     Journal
LA
     English
     28-23 (Heterocyclic Compounds (More Than One Hetero Atom))
CC
     Section cross-reference(s): 68
os
     CASREACT 135:303868
GI
* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *
     Calix[4] arene derivs. such as I and II (R = EtCH2S)
     contq. N or S atoms at the lower rim were prepd. E.g., di(2-bromoethyl)
     calixarene II (R = Br) was refluxed with 1,3-propanedithiol in a
     2:1 mixt. of benzene and DMF for 4h with potassium hydroxide as the base
     to give I in 87% yield. All of the receptors prepd. were highly selective
     for silver ion over other metal ions. The calix[4] arene podand
     II (R = EtCH2S) bound Aq+ more strongly than the calix
     [4] crown ligands such as I.
ST
     calixcrown calixarene podand ligand prepn selective
     extn silver; complexation extn metal ion calixcrown
     calixarene podand ligand; selective complexation silver
     calixcrown calixarene podand
     Metacyclophanes
TΤ
     RL: PRP (Properties); RCT (Reactant); SPN (Synthetic preparation); PREP
     (Preparation); RACT (Reactant or reagent)
        (calixarenes; prepn. and metal ion complexation selectivity
        of calixarene thiacrown and podand derivs. as ligands for the
        selective complexation of silver)
     Metacyclophanes
IT
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (crown ethers; prepn. and metal ion complexation selectivity of
        calixarene thiacrown and podand derivs. as ligands for the
        selective complexation of silver)
     Crown ethers
IT
```

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (metacyclophanes; prepn. and metal ion complexation selectivity of calixarene thiacrown and podand derivs. as ligands for the selective complexation of silver) Complexation (prepn. and metal ion complexation selectivity of calixarene thiacrown and podand derivs. as ligands for the selective complexation of silver) Macrocyclic compounds RL: PRP (Properties) (prepn. and metal ion complexation selectivity of calixarene thiacrown and podand derivs. as ligands for the selective complexation of silver) 7439-89-6, Iron, properties 7439-96-5, Manganese, properties 7440-09-7, Potassium, properties 7439-97-6, Mercury, properties 7440-22-4, Silver, properties 7440-28-0, Thallium, properties 7440-43-9, Cadmium, properties 7440-48-4, Cobalt, properties 7440-50-8, Copper, properties 7440-66-6, Zinc, properties 7440-70-2, Calcium, properties 223911-14-6 223911-20-4 223911-21-5 RL: PRP (Properties) (prepn. and metal ion complexation selectivity of calixarene thiacrown and podand derivs. as ligands for the selective complexation of silver) 197228-56-1P 367279-78-5P 367279-79-6P RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (prepn. and metal ion complexation selectivity of calixarene thiacrown and podand derivs. as ligands for the selective complexation of silver) 107-03-9, 1-Propanethiol 109-64-8, 1,3-Dibromopropane 109-80-8, 1,3-Propanedithiol 60705-62-6 197228-58-3 RL: RCT (Reactant); RACT (Reactant or reagent) (prepn. and metal ion complexation selectivity of calixarene thiacrown and podand derivs. as ligands for the selective complexation of silver) 213924-21-1P RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (prepn. and metal ion complexation selectivity of calixarene thiacrown and podand derivs. as ligands for the selective complexation of silver) RE.CNT THERE ARE 29 CITED REFERENCES AVAILABLE FOR THIS RECORD (1) Aggarwal, R; Def Sci J 1975, V25, P153 CA (2) Bakker, W; J Chem Soc, Chem Commun 1994, P71 CA (3) Bohmer, V; Angew Chem, Int Ed 1995, V34, P715 (4) Casnati, A; Chem Eur J 1996, V2, P436 CA (5) Casnati, A; J Am Chem Soc 1995, V117, P2767 CA (6) Chen, C; Synth Commun in press (7) Couton, D; Aust J Chem 1999, V52, P227 CA (8) Covington, A; J Chem Thermodynamics 1971, V3, P795 CA (9) Dijkstra, P; J Am Chem Soc 1989, V111, P7567 CA (10) Ghidini, E; J Am Chem Soc 1990, V112, P6979 CA (11) Gutsche, C; Calixarene 1989 (12) Gutsche, C; Calixarene Revisited 1998 (13) Gutsche, C; Org Synth 1989, V68, P234 (14) Harris, S; EP 0432989 A2 1990 CA (15) Ikeda, A; Chem Rev 1997, V97, P1713 CA (16) Inoue, Y; Bull Chem Soc 1985, V58, P525 CA

IT

IT

IT

IT

IT

IT

RE

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   P127 CA
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(28) Zhenq, Q; J Chem Res S 1999, P212 CA
(29) Zheng, Y; Chinese Chem Lett 1997, V8, P685 CA
L3
     ANSWER 3 OF 30 CA COPYRIGHT 2002 ACS
     135:203004 CA
AN
ΤI
     Post-development resist hardening by vapor silylation
     Aviram, Ari; Rooks, Michael Joseph
IN
PA
     International Business Machines Corp., USA
so
     U.S., 7 pp.
     CODEN: USXXAM
DT
     Patent
LΑ
     English
IC
     ICM G03F007-26
NCL
    430314000
     74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other
CC
     Reprographic Processes)
     Section cross-reference(s): 76
FAN.CNT 1
     PATENT NO.
                    KIND DATE
                                         APPLICATION NO. DATE
     -----
                                          -----
                     B1 20010828
                                         US 1999-292727 19990415
PT
     The invention provides a method of reducing the etch rate of a
AB
     patterned resist on a substrate by treating the patterned resist with an
     atm. comprising the mols. of a hardening agent. Specifically, the method
     employed by the present invention comprises applying a layer of an
     imageable resist to a substrate layer; patterning the layer of imageable
     resist by removing selective areas thereof; and treating the patterned
     imageable resist with an atm. comprising mols. of a hardening agent so as
     to obtain a hardened resist surface which etches at a slower
     rate than that of the untreated resist.
     resist hardening vapor silylation methyl acetoxy calixarene
st
     hexamethyl disilazane
IT
     Etching kinetics
        (in CF4 plasma, of methylacetoxycalixarene resist film hardened by
        vapor silylation using hexamethyldisilazane)
IT
     Resists
        (pos.-tone; post-development resist hardening of
        methylacetoxycalixarene by vapor silylation using hexamethyldisilazane)
IT
     Silvlation
     Surface hardening
        (post-development resist hardening of methylacetoxycalixarene by vapor
        silylation using hexamethyldisilazane)
IT
     141137-71-5
     RL: DEV (Device component use); PEP (Physical, engineering or chemical
     process); TEM (Technical or engineered material use); PROC (Process); USES
        (post-development resist hardening by vapor silylation of)
IT
     999-97-3, HMDS
     RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical
     process); RCT (Reactant); PROC (Process); RACT (Reactant or reagent); USES
     (Uses)
        (post-development resist hardening of methylacetoxycalixarene by vapor
        silylation using)
IT
     7439-88-5, Iridium, uses 7439-89-6, Iron, uses 7439-95-4, Magnesium,
           7440-02-0, Nickel, uses 7440-05-3, Palladium, uses 7440-06-4,
     Platinum, uses 7440-16-6, Rhodium, uses 7440-18-8, Ruthenium, uses
     7440-32-6, Titanium, uses 7440-38-2, Arsenic, uses 7440-48-4, Cobalt,
           7440-50-8, Copper, uses 7440-57-5, Gold, uses 7440-67-7,
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Zirconium, uses 7440-70-2, Calcium, uses RL: DEV (Device component use); MOA (Modifier or additive use); USES (post-development resist hardening of methylacetoxycalixarene using metalloid or metallic element) RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD (1) Bottcher, M; J Vac Sci Technol B 1994, V12(6), P3473 (2) Hien; US 6063543 2000 CA (3) Irmscher, M; J Vac Sci Technol B 1994, V12(6), P3925 CA (4) McCord, M; "Chapter 2 Electron Beam Lithography," Handbook of Microlithography, Micromachining and Microfabrication vol 1: Microlithography 1997 (5) Sato; US 5756255 1998 CA ANSWER 4 OF 30 CA COPYRIGHT 2002 ACS 135:53439 CA Microcontact printing with heavyweight inks Liebau, Maik; Huskens, Jurriaan; Reinhoudt, David N. Laboratory of Supramolecular Chemistry and Technology, MESA+ Research Institute, University of Twente, Enschede, NL-7500 AE, Neth. Advanced Functional Materials (2001), 11(2), 147-150 CODEN: AFMDC6; ISSN: 1616-301X Wiley-VCH Verlag GmbH Journal English 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes) Thioether derivs. with thioether moieties were used as inks in microcontact printing on gold for the reprodn. of patterns as they combine good monolayer quality with synthetic versatility, and high mol. wts. Self-assembled monolayers (SAMs) on gold of calixarene- and cavitand tetra(thioether) compds. (1 and 2 resp.) had a quality comparable to SAMs of decanethiol, both regarding monolayer order and etch resistance. Etch resistances of SAMs of tris(thioether) deriv. and dodecylsulfide (3 and 4 resp.) were lower. Resulting structures after etching of patterned SAMs using 1, 2, and 3 were of good quality for patterns with feature sizes on the same stamp ranging from 1-200 .mu.m. Patterns of 4 were not reproduced. Overall, compds. 1 and 2 are good candidates for low-diffusion inks. lithog microcontact printing thioether deriv self assembled layer; calixarene thioether deriv self assembled layer lithog microcontact printing Metacyclophanes RL: PEP (Physical, engineering or chemical process); PROC (Process) (calixarenes; etch resistance and imaging properties of self-assembled monolayers of thioether derivs. in microcontact printing of gold surfaces) Electric capacitance Electric resistance (electrochem. - and imaging properties of self-assembled monolayers of thioether derivs. in microcontact printing of gold surfaces) Contact angle Self-assembled monolayers (etch resistance and imaging properties of self-assembled monolayers of thioether derivs. in microcontact printing of gold surfaces) Thioethers RL: PEP (Physical, engineering or chemical process); PROC (Process) (etch resistance and imaging properties of self-assembled monolayers of thioether derivs. in microcontact printing of gold

IT Lithography (microcontact printing; etch resistance and imaging

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surfaces)

properties of self-assembled monolayers of thioether derivs. in microcontact printing of gold surfaces) IT 693-83-4, Didecyl sulfide RL: PEP (Physical, engineering or chemical process); PRP (Properties); PROC (Process) (comparison; etch resistance and imaging properties of self-assembled monolayers of thioether derivs. in microcontact printing of gold surfaces) 7440-57-5, Gold, processes IT RL: PEP (Physical, engineering or chemical process); PROC (Process) (etch resistance and imaging properties of self-assembled monolayers of thioether derivs. in microcontact printing of gold surfaces) IT 155401-92-6 167424-00-2 345196-83-0 RL: PEP (Physical, engineering or chemical process); PRP (Properties); PROC (Process) (etch resistance and imaging properties of self-assembled monolayers of thioether derivs. in microcontact printing of gold surfaces) IT 1310-58-3, Potassium hydroxide, uses 7772-98-7, Sodium thiosulfate 13943-58-3, Tetrapotassium hexacyanoferrate RL: NUU (Other use, unclassified); USES (Uses) (etchant; etch resistance and imaging properties of self-assembled monolayers of thioether derivs. in microcontact printing of gold surfaces) RE.CNT THERE ARE 17 CITED REFERENCES AVAILABLE FOR THIS RECORD 17 RE(1) Beulen, M; Langmuir 1996, V12, P6170 CA (2) Boukamp, B; Solid State Ionics 1986, V20, P31 CA (3) Delamarche, E; J Phys Chem B 1998, V102, P3324 CA (4) Dubois, H; J Chem Phys 1993, V98, P678 (5) Eberhardt, A; Langmuir 1999, V15, P1595 CA (6) Huangxian, J; Langmuir 1998, V14, P300 (7) Huisman, B; Tetrahedron Lett 1995, V36, P3273 CA (8) Larsen, B; J Am Chem Soc 1997, V119, P3017 (9) Li, X; unpublished (10) Libiuouelle, L; Langmuir 1999, V15, P300 (11) Schmid, H; Macromolecules 2000, V33, P3042 CA (12) Schonherr, H; Langmuir 1999, V15, P5541 (13) van Velzen, E; J Am Chem Soc 1994, V116, P3597 (14) van Velzen, E; J Am Chem Soc 1995, V117, P6853 (15) Xia, Y; Angew Chem Int Ed 1998, V37, P550 CA (16) Xia, Y; Chem Mater 1995, V7, P2332 CA (17) Zhao, X; Langmuir 1996, V12, P3257 CA L3 ANSWER 5 OF 30 CA COPYRIGHT 2002 ACS AN 134:229590 CA ΤI Comparative study of sputtered and spin-coatable aluminum oxide electron beam resists AU Saifullah, M. S. M.; Kurihara, K.; Humphreys, C. J. Nanoscale Science Group, Department of Engineering, University of CS Cambridge, Cambridge, CB2 1PZ, UK so Journal of Vacuum Science & Technology, B: Microelectronics and Nanometer Structures (2000), 18(6), 2737-2744 CODEN: JVTBD9; ISSN: 0734-211X PB American Institute of Physics DT Journal LA English CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes) AB The electron beam exposure characteristics of sputtered AlOx and spin-coatable Al203 resists are compared and contrasted. When exposed to an electron beam, sputtered AlOx resists on a silicon substrate undergo an

intense mass loss. However, electron energy loss spectroscopy shows that

even after a prolonged exposure some aluminum and oxygen remains on the silicon surface. Spin-coatable Al2O3 resist was prepd. by reacting aluminum tri-sec-butoxide, Al(OBus)3, with acetylacetone (AcAc) in iso-Pr These are neg. tone resists and they are >106 times more sensitive to an electron beam than the sputtered AlOx, bringing its sensitivity very close to high resoln. org. resists such as calixarene. The exposure properties of spin-coatable and sputtered aluminum oxide resists are discussed together with their sensitivity, damage mechanisms, line edge roughness, and etching characteristics. A brief note on the change of methodol. of resist design is added when inorg. resists are to be used in high resoln. electron beam nanolithog. sputtered spin coated aluminum oxide lithog electron beam resist Electron beam resists Electron radiolysis (comparison of lithog. characteristics of sputtered and spin-coated aluminum oxide electron-beam resists) Electron beams IR spectra (lithog. characteristics of spin coated aluminum oxide electron-beam resist prepd. from reaction of aluminum butoxide with acetylacetone) 123-54-6, Acetylacetone, uses RL: NUU (Other use, unclassified); USES (Uses) (comparison of lithog. characteristics of sputtered and spin-coated aluminum oxide electron-beam resists) 1344-28-1, Aluminum oxide, properties RL: PEP (Physical, engineering or chemical process); PRP (Properties); TEM (Technical or engineered material use); PROC (Process); USES (Uses) (comparison of lithog. characteristics of sputtered and spin-coated aluminum oxide electron-beam resists) 2269-22-9, Aluminum tri-sec-butoxide RL: NUU (Other use, unclassified); USES (Uses) (lithog. characteristics of spin coated aluminum oxide electron-beam resist prepd. from reaction of aluminum butoxide with acetylacetone) THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT (1) Berger, S; Philos Mag B 1987, V55, P341 CA (2) Chen, G; Appl Phys Lett 1993, V62, P1949 CA (3) Chen, W; Appl Phys Lett 1993, V62, P1499 CA (4) Chen, W; J Vac Sci Technol B 1993, V11, P2519 CA (5) Devenish, R; Ultramicroscopy 1989, V28, P324 (6) Fujita, J; Appl Phys Lett 1995, V66, P3065 CA (7) Hollenbach, J; J Mater Res 1990, V5, P1058 (8) Knotek, M; Phys Rev Lett 1978, V40, P964 CA (9) Kratschmer, E; J Vac Sci Technol B 1987, V5, P369 CA

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L3
AN
     134:23509 CA
    Method for pattern formation using calix[7] arene for
TI
     semiconductor substrate
     Shinko, Sachiko; Ochiai, Yukinori; Yamamoto, Hiromasa; Tejima, Takahiro
IN
    NEC Corp., Japan; Tokuyama Corp.
PΑ
     Jpn. Kokai Tokkyo Koho, 7 pp.
SO
    CODEN: JKXXAF
DT
    Patent
LA
     Japanese
     ICM G03F007-038
IC
     ICS H01L021-027
     74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other
CC
     Reprographic Processes)
     Section cross-reference(s): 76
FAN.CNT 1
                                         APPLICATION NO. DATE
                    KIND DATE
    PATENT NO.
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     JP 2000330281 A2 20001130
                                          JP 1999-144369 19990525
ΡI
     The title method includes the steps of: (1) forming a thin layer on a
AΒ
     substrate using a coating soln. mainly made of calix[7]arene;
     (2) forming a latent image on the thin layer using a high energy beam; and
     (3) selectively etching-off the thin layer except the latent
     image part to form a pattern. The method using calix[7] arene
     provides a pattern of the reduced pattern roughness.
ST
     pattern formation resist semiconductor substrate calixarene
IT
     Metacyclophanes
     RL: TEM (Technical or engineered material use); USES (Uses)
        (calixarenes; light sensitive compn. for semiconductor
        substrate prodn.)
ΙT
     Photoresists
     Semiconductor device fabrication
        (method for pattern formation for semiconductor substrate)
     50-00-0, Formaldehyde, reactions 106-44-5, p-Cresol, reactions
ΙT
     108-24-7, Acetic anhydride
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (calixarene in light-sensitive resist compn.)
     141137-71-5P, p-Methylhexaacetoxycalix[6]arene
                                                     196408-88-5P,
     p-Methylheptaacetoxycalix[7] arene 196408-89-6P, p-
     Methyloctaacetoxycalix[8] arene
     RL: SPN (Synthetic preparation); TEM (Technical or engineered material
     use); PREP (Preparation); USES (Uses)
        (calixarene in light-sensitive resist compn.)
     ANSWER 7 OF 30 CA COPYRIGHT 2002 ACS
L3
AN
     A comparison between the complexant alkaline etching systems of
TI
ΑU
     Moldovan, Carmen; Iosub, Rodica; Dascalu, Dan; Mrin, Gheorghe; Danila,
     Natl. Inst. R&D Microtechnol., Bucharest, Rom.
CS
     CAS '99 Proceedings, International Semiconductor Conference, 22nd, Sinaia,
SO
     Romania, Oct. 5-9, 1999 (1999), Volume 1, 205-208 Publisher: Institute of
     Electrical and Electronics Engineers, New York, N. Y.
     CODEN: 69ADUW
DT
     Conference
LΑ
     English
     76-3 (Electric Phenomena)
CC
     This paper contains the results obtained in silicon hillock elimination
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using alk. solns.: KOH, NaOH, LiOH - H2O with complexant added. complexant added in alk. solns. is Azo calix[4] arene. The alk.

solns. were compared and analyzed with and without complexant added, in point of view of hillocks and the behavior of these solns. is explained using the theory of molar cond. The results allow the authors to use the alk. solns. and the org. complexant, to monitor the etching process, to obtain a smooth silicon surface, almost free of hillocks. alk etching systems silicon surface; alkali metal hydroxide etching system silicon surface Semiconductor materials Surface structure (a comparison between the complexant alk. etching systems of silicon) Alkali metal hydroxides RL: RCT (Reactant); RACT (Reactant or reagent) (a comparison between the complexant alk. etching systems of silicon) 7440-21-3P, Silicon, properties RL: DEV (Device component use); PNU (Preparation, unclassified); PRP (Properties); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses) (a comparison between the complexant alk. etching systems of silicon) 221181-37-9D, salts RL: FMU (Formation, unclassified); PRP (Properties); RCT (Reactant); FORM (Formation, nonpreparative); RACT (Reactant or reagent) (a comparison between the complexant alk. etching systems of silicon) 1310-58-3, Potassium hydroxide (KOH), reactions 1310-65-2, Lithium 1310-73-2, Sodium hydroxide (NaOH), reactions hydroxide (LiOH) RL: RCT (Reactant); RACT (Reactant or reagent) (a comparison between the complexant alk. etching systems of silicon) THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT 5 (1) Atkins, P; Physical Chemistry 1994, P765 (2) Moldovan, C; EUROSENSORS XII V2, P1009 CA (3) Moldovan, C; Sensors and Actuators B 1999, V000, P1 (4) Moldovan, C; proceedings of SPIE, Design, Test and Microfabrication of MEMS and MOEMS 1999, P1056 CA (5) Seidel, H; JES 1990, V137, P3612 CA ANSWER 8 OF 30 CA COPYRIGHT 2002 ACS 133:244989 CA Nanometer-scale resolution of a chloromethylated calixarene negative resist in electron-beam lithography: Dependence on the number of phenolic residues Sakamoto, T.; Manako, S.; Fujita, J.; Ochiai, Y.; Baba, T.; Yamamoto, H.; Teshima, T. Fundamental Research Laboratories, NEC Corporation, Tsukuba, Ibaraki, 305-8501, Japan Applied Physics Letters (2000), 77(2), 301-303 CODEN: APPLAB; ISSN: 0003-6951 American Institute of Physics Journal English 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes) We have investigated a chloromethylated calixarene, p-chloromethylmethoxcalix[n]arene (CMC[n]AOMe) (n=5,6,7), as a neg. resist in electron-beam lithog. Each CMC[n] AOMe resist has a resoln. of about 12 nm and a sensitivity of about 0.8 mC/cm2 which varies slightly with n (or mol. wt.). A sub-10-nm Si wire has been fabricated by halide plasma etching and a CMC(n) AOMe resist as an etching mask. Because the resist pattern edge is smooth, Si wires with 7-nm width and 10-.mu.m length were performed without any breaking.

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AB

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ST
     chloromethylated calixarene electron beam resist; lithog
     electron beam etching mask
IT
     Electron beam lithography
     Electron beam resists
       Etching
       Etching masks
        (nanometer-scale resoln. of chloromethylated calixarene neg.
        resist in electron-beam lithog.: Dependence on no. of phenolic
        residues)
     7440-21-3, Silicon, processes
IT
     RL: DEV (Device component use); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (nanometer-scale resoln. of chloromethylated calixarene neg.
        resist in electron-beam lithog.: Dependence on no. of phenolic
        residues)
RE.CNT 12
              THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
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L3
     ANSWER 9 OF 30 CA COPYRIGHT 2002 ACS
AN
     133:176912 CA
ΤI
     conformational rigidity in mesitylene-based calix[4] arenes
     adopting a 1,3-alternate conformation
ΑU
     Parzuchowski, P.; Bohmer, V.; Biali, S. E.; Thondorf, I.
CS
     ul. Noakowskiego 3, Faculty of Chemistry, Warsaw University of Technology,
     Warsaw, PL-00664, Pol.
SO
     Tetrahedron: Asymmetry (2000), 11(11), 2393-2402
     CODEN: TASYE3; ISSN: 0957-4166
PB
     Elsevier Science Ltd.
DT
     Journal
LΑ
     English
CC
     22-10 (Physical Organic Chemistry)
GI
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Two chiral derivs. of a mesitylene-based calix[4] arene (I; R=
EtCHMeCH2, PhCHMeNHCOCH2) known to exist in the 1,3-alternate
conformation were prepd. by the attachment of homochiral residues to the
four exo-hydroxy groups. Thus, the enantiotopic protons of the central
scaffold became diastereotopic, leading to a doubling of their 1H NMR
signals in one example. From the temp. independence of the NMR spectrum,
a lower limit of 24.2 kcal/mol could be estd. for the barrier of ring
inversion. MM3 calcns. confirm the 1,3-alternate conformation as the
energy min., and est. a barrier of 25.7 kcal/mol for the
1,3-alternate-to-1,3-alternate* interconversion process. This high
barrier is due to the repulsive steric interactions between exo-Me groups
at vicinal rings when these groups pass each other.

ST NMR conformation calixarene chiral mesitylene based; rotational barrier calixarene chiral mesitylene based NMR

IT Steric hindrance

Strain energy

(Me group; NMR study of conformational rigidity in mesitylene-based calix[4] arenes adopting a 1,3-alternate conformation)

IT Chirality

Conformation

Conformational potential

Conformational transition

Molecular mechanics

NMR (nuclear magnetic resonance)

Rotational barrier

(NMR study of conformational rigidity in mesitylene-based calix [4] arenes adopting a 1,3-alternate conformation)

IT Metacyclophanes

RL: PEP (Physical, engineering or chemical process); PRP (Properties); PROC (Process)

(calixarenes; NMR study of conformational rigidity in mesitylene-based calix[4] arenes adopting a 1,3-alternate conformation)

IT Conformers

(low energy; NMR study of conformational rigidity in mesitylene-based calix[4] arenes adopting a 1,3-alternate conformation)

IT Methyl group

(steric interaction; NMR study of conformational rigidity in mesitylene-based calix[4] arenes adopting a 1,3-alternate conformation)

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IT
                   149703-31-1
                                 157432-87-6
                                               288301-95-1
                                                             288302-11-4
     121702-02-1
     288302-12-5 288302-13-6
                                 288302-14-7
                                               288302-15-8
                                                             288302-16-9
     288302-17-0
                   288302-18-1
     RL: PEP (Physical, engineering or chemical process); PRP (Properties);
     PROC (Process)
        (NMR study of conformational rigidity in mesitylene-based calix
        [4] arenes adopting a 1,3-alternate conformation)
IT
     288301-93-9P
                    288301-94-0P
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); SPN
     (Synthetic preparation); PREP (Preparation); PROC (Process)
        (NMR study of conformational rigidity in mesitylene-based calix
        [4] arenes adopting a 1,3-alternate conformation)
IT
     534-00-9, (S)-(+)-1-Bromo-2-methylbutane
                                                2627-86-3, (S)-(-)-1-
     Phenylethylamine
                        201489-72-7
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (prepn. of chiral derivs. of a mesitylene-based calix
        [4] arene)
IT
     212835-17-1P
     RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
     (Reactant or reagent)
        (prepn. of chiral derivs. of a mesitylene-based calix
        [4]arene)
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L3
     ANSWER 10 OF 30 CA COPYRIGHT 2002 ACS
AN
     133:170174 CA
ΤI
     Electron beam lithography of Fresnel zone plates using a rectilinear
     machine and trilayer resists
ΑU
     Tennant, D.; Spector, S.; Stein, A.; Jacobsen, C.
     Lucent Technologies Bell Laboratories, USA
CS
SO
     AIP Conference Proceedings (2000), 507(X-Ray Microscopy), 601-606
     CODEN: APCPCS; ISSN: 0094-243X
PΒ
     American Institute of Physics
DT
     Journal
LA
     English
CC
     74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other
     Reprographic Processes)
     Section cross-reference(s): 73
AB
     The authors describe the use of a com. e-beam lithog. system (JEOL
     JBX-6000FS) to fabricate Fresnel zone plates for x-ray microscopy.
     machine is capable of controlling the pitch of optical gratings with
     sub-nanometer precision, so its beam placement properties are more than
     adequate for zone plate fabrication. The zone plate pattern is written
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into a thin top layer (PMMA or Calixarene) of a trilayer resist,

and transferred into thick nickel zones using reactive ion **etching** (RIE) followed by electroplating. Zone plates with outermost zone widths of 30 nm have exhibited efficiencies up to 10.0% at a 390 eV photon energy and with diams. in the range 80 to 120 .mu.m. Zone plates with outer zones of 18 to 20 nm were also fabricated in thinner Ni with correspondingly lower efficiencies of 2.6%. Zone plates with outermost zone widths of 45 nm have been fabricated with larger diams. up to 160 .mu.m. All results reported were obtained with a 50 kV system with 80 .mu.m field deflection size; future efforts will make use of a 100 kV, 500 .mu.m field size system.

ST Fresnel zone plate fabrication electron beam lithog

IT Metacyclophanes

RL: TEM (Technical or engineered material use); USES (Uses)
(calixarenes; electron beam lithog. fabrication of Fresnel
zone plates for x-ray microscopy using rectilinear machine and trilayer
resists)

IT Electron beam lithography

Electron beam resists

X-ray microscopy

(electron beam lithog. fabrication of Fresnel zone plates for x-ray microscopy using rectilinear machine and trilayer resists)

IT Optical instruments

(zone plates, Fresnel; electron beam lithog. fabrication of Fresnel zone plates for x-ray microscopy using rectilinear machine and trilayer resists)

IT 89072-52-6, AZ 4110

RL: NUU (Other use, unclassified); USES (Uses) (AZ 4110; electron beam lithog. fabrication of Fresnel zone plates for x-ray microscopy using rectilinear machine and trilayer resists)

IT 7440-02-0, Nickel, processes 7440-47-3, Chrome, processes 7440-57-5 Gold, processes

RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)

(electron beam lithog. fabrication of Fresnel zone plates for x-ray microscopy using rectilinear machine and trilayer resists)

IT 7440-56-4, Germanium, uses 9011-14-7, PMMA

RL: NUU (Other use, unclassified); USES (Uses)
(electron beam lithog. fabrication of Fresnel zone plates for x-ray
microscopy using rectilinear machine and trilayer resists)

IT 75-25-2, Tribromomethane

RL: NUU (Other use, unclassified); USES (Uses)
 (plasma etch; electron beam lithog. fabrication of Fresnel
 zone plates for x-ray microscopy using rectilinear machine and trilayer
 resists)

RE.CNT 13 THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS RECORD RE

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- L3 ANSWER 11 OF 30 CA COPYRIGHT 2002 ACS

AN 133:134995 CA

TI Non-directional forces drive guest affinity and discrimination in a calix[5] arene-based receptor

AU Haino, Takeharu; Nitta, Koji; Fukazawa, Yoshimasa

CS Department of Chemistry, Graduate School of Science, Hiroshima University, Higashi-Hiroshima, 739-8526, Japan

SO Tetrahedron Letters (2000), 41(21), 4139-4142 CODEN: TELEAY; ISSN: 0040-4039

PB Elsevier Science Ltd.

DT Journal

LA English

CC 22-10 (Physical Organic Chemistry)

GI

AB In this paper we report the binding behavior of a receptor based on a calix[5] arene possessing two convergent benzoic acids that serve to bind a guest. The receptor recognizes diketopiperazines (DKP) having a variety of alkyl substituents to form a 1:1 host-guest complex (I; R= Me,Me2CH,EtCHMe,Me2CHCH2,Me3C,MeSCH2CH2, PhCH2). We demonstrate that the receptor shows selectivity for the DKPs based on the size of the alkyl substituent. As a result, the selectivity arises from non-directional forces between the .pi.-basic arom. ring of the calixarene and the alkyl group of the guest.

Ι

ST calixarene receptor diketopiperazine guest affinity NMR mol mechanics; assocn const calixarene receptor diketopiperazine quest

IT Formation constant

Hydrogen bond

Molecular mechanics

Molecular modeling

NMR (nuclear magnetic resonance)

Van der Waals force

(NMR and mol. mechanics study of guest affinity and discrimination in calix[5]arene-based receptor)

IT Molar volume

Molecular shape

(alkyl group; NMR and mol. mechanics study of guest affinity and discrimination in calix[5] arene-based receptor)

IT Metacyclophanes

RL: PEP (Physical, engineering or chemical process); PROC (Process) (calixarenes; NMR and mol. mechanics study of guest affinity and discrimination in calix[5] arene-based receptor)

```
TΤ
     Solvation
        (model; NMR and mol. mechanics study of guest affinity and
        discrimination in calix[5] arene-based receptor)
IT
     Force
        (nondirectional; NMR and mol. mechanics study of guest affinity and
        discrimination in calix[5] arene-based receptor)
IT
     Alkyl groups
        (shape anal.; NMR and mol. mechanics study of guest affinity and
        discrimination in calix[5] arene-based receptor)
                    286856-04-0P
                                   286856-05-1P
                                                  286856-06-2P
                                                                  286856-08-4P
ΙT
     286856-03-9P
                    286856-11-9P
     286856-10-8P
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
        (NMR and mol. mechanics study of guest affinity and discrimination in
        calix(5) arene-based receptor)
RE.CNT
              THERE ARE 23 CITED REFERENCES AVAILABLE FOR THIS RECORD
       23
RE
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(18) Nicholls, A; Proteins 1991, V11, P281 CA
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(20) Rebek, J; Acc Chem Res 1990, V23, P399 CA
(21) Shao, Y; J Org Chem 1996, V61, P6086 CA
(22) Szabo, T; Angew Chem, Int Ed 1998, V37, P3410 CA
(23) Yoon, S; Tetrahedron 1995, V51, P567 CA
L3
     ANSWER 12 OF 30 CA COPYRIGHT 2002 ACS
AN
     132:257193 CA
TI
     Aspects concerning chemical and electrochemical reactivity of some
     synthesized calixarenes
AU
     Unqureanu, Eleonora-Mihaela; Nechifor, G.; Pirvu, C.; Serban, B.
     Department of Industrial Chemistry, University "Politehnica" of Bucharest,
CS
     Scientific Bulletin - University "Politehnica" of Bucharest, Series B:
so
     Chemistry and Materials Science (1999), 61(1-2), 55-65
     CODEN: SBUPBD; ISSN: 1454-2331
PΒ
     University "Politehnica" of Bucharest
DT
     Journal
LA
     English
     72-2 (Electrochemistry)
CC
     Section cross-reference(s): 22, 25, 66, 76
     Study of electrochem. properties of newly synthesized calix
AB
     [4] arenes and the silicon etching in basic medium in the
     presence of these calixarenes are described.
ST
     electrochem chem reactivity calixarene prepn; silicon
     anisotropic etching alk soln calixarene complexant;
     electroredn calixarene glassy carbon platinum electrode;
     electrooxidn calixarene glassy carbon platinum electrode; cyclic
     voltammetry calixarene glassy carbon platinum electrode
IT
     Etching
```

```
(anisotropic; of silicon in alk. soln. contq. calixarene
        sulfophenylazo deriv. complexant)
IT
     Reactivity (chemical)
        (chem. and electrochem. reactivity of synthesized calixarenes
ΙT
     Cyclic voltammetry
     Oxidation, electrochemical
     Reduction, electrochemical
        (of calixarenes on glassy carbon or platinum in acetonitrile)
ΙT
     UV and visible spectra
        (of sulfophenylazo deriv. of calixarene)
IT
     Substituent effects
        (on cyclic voltammetry of calixarenes)
IT
     7440-21-3, Silicon, properties
     RL: PEP (Physical, engineering or chemical process); PRP (Properties);
     PROC (Process)
        (anisotropic etching of silicon in alk. soln. contg.
        calixarene complexant)
ΙT
     124411-21-8
     RL: NUU (Other use, unclassified); PRP (Properties); RCT (Reactant); RACT
     (Reactant or reagent); USES (Uses)
        (anisotropic etching of silicon in alk. soln. contg.
        complexant)
ΙT
     7440-06-4, Platinum, uses
                                 7440-44-0, Carbon, uses
     RL: DEV (Device component use); PRP (Properties); USES (Uses)
        (cyclic voltammetry of calixarenes on glassy carbon or
        platinum in acetonitrile)
IT
     429-42-5, Tetrabutylammonium tetrafluoroborate
     RL: NUU (Other use, unclassified); PRP (Properties); USES (Uses)
        (cyclic voltammetry of calixarenes on glassy carbon or
        platinum in acetonitrile contg.)
     7446-70-0, Aluminum chloride, uses
IT
     RL: CAT (Catalyst use); USES (Uses)
        (in debutylation of calixarene tert-Bu deriv. in phenol)
IT
     74568-07-3P
                   123796-82-7P
                                  262428-23-9P
                                                  262428-24-0P
     RL: PRP (Properties); RCT (Reactant); SPN (Synthetic preparation); PREP
     (Preparation); RACT (Reactant or reagent)
        (prepn. and cyclic voltammetry on glassy carbon or platinum in
        acetonitrile)
IT
     60705-62-6P
     RL: PRP (Properties); RCT (Reactant); SPN (Synthetic preparation); PREP
     (Preparation); RACT (Reactant or reagent)
        (prepn. and dealkylation in toluene contg. phenol and aluminum chloride
        and cyclic voltammetry on platinum in acetonitrile)
TT
     100-05-0
                119-66-4
                           262428-25-1
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reaction with calixarene in NaOH)
IT
     98-54-4, p-tert-Butylphenol
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reaction with formaldehyde in NaOH in calixarene tert-Bu
        deriv. prepn.)
TT
     50-00-0, Formaldehyde, reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reaction with p-tert-butylphenol in NaOH in calixarene
        tert-Bu deriv. prepn.)
RE.CNT 20
              THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
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(2) Beer, P; Molecular Engineering for Advanced Materials 1989
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(9) Moldovan, C; EUROSENSORS XII 1998
(10) Moldovan, C; Workshop on Physical Chemistry of Wet Chemical Etching of
    Silicon 1998
(11) Moutet, J; J Electroanal Chem 1996, V410, P79 CA
(12) Ristic, L; Sensor Technology and Devices 1994, P67
(13) Seidel, H; J Electrochem Soc 1990, V137(11), P3612 CA
(14) Shinkai, S; J Am Chem Soc 1986, V108, P2409 CA
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(20) Verboon, W; Molecular Engineering for Advanced Materials 1989
L3
     ANSWER 13 OF 30 CA COPYRIGHT 2002 ACS
ΑN
     132:71961 CA
TI
     Anisotropic etching of silicon in a complexant redox alkaline
ΑU
     Moldovan, Carmen; Iosub, Rodica; Dascalu, Dan; Nechifor, Gheorghe
CS
     National Institute for R and D in Microtechnologies, Bucharest, R72225,
     Rom.
SO
     Sensors and Actuators, B: Chemical (1999), B58(1-3), 438-449
     CODEN: SABCEB; ISSN: 0925-4005
PB
     Elsevier Science S.A.
DT
     Journal
LΑ
     English
CC
     76-3 (Electric Phenomena)
AB
     This paper presents the results from the investigation of the chem.
     anisotropic etching of single-crystal silicon 100 in the
     following solns.: KOH, K3[Fe(CN)6] 0.1 M, K4[Fe(CN)6].cntdot.3H2O 0.1 M,
     KNO3 0.1 M and or complexant added. The complexant added in KOH soln.
     were: calix[4] arenes, phenols and ether dibenzo-18-crown-6.
     reaction mechanism, the etch rate, the roughness and the
     hillocks are analyzed. The results allow us to use the redox system
     and/or the org. complexants to monitor the etching process, to
     obtain a smooth silicon surface with increased etch rate and to
     utilize the usual mask material resistant at the new etchants.
st
     silicon anisotropic etching etchant redox alk
ΙT
     Surface roughness
        (after etching anal. of; anisotropic etching of
        silicon in a complexant redox alk. system)
IT
        (anisotropic, of silicon; anisotropic etching of silicon in a
        complexant redox alk. system)
IT
     Redox reaction
        (complexant, alk. system; anisotropic etching of silicon in a
        complexant redox alk. system)
IT
     Surface structure
        (hillocks, after etching anal. of; anisotropic
        etching of silicon in a complexant redox alk. system)
IT
     98-54-4, p-tert-Butylphenol
                                  108-95-2, Phenol, uses
                     14187-32-7, Dibenzo-18-crown-6
     Calix[4] arene
                                                      25154-52-3,
     Nonylphenol
                   60705-62-6, p-tert-Butyl calix[4] arene
     RL: MOA (Modifier or additive use); USES (Uses)
        (anisotropic etching of silicon in a complexant redox alk.
        system)
     7440-21-3, Silicon, properties
IT
     RL: PEP (Physical, engineering or chemical process); PRP (Properties);
     PROC (Process)
        (anisotropic etching of silicon in a complexant redox alk.
        system)
IT
     221181-37-9
    RL: MOA (Modifier or additive use); USES (Uses)
```

(azo calix[4]arene; anisotropic etching of silicon
in a complexant redox alk. system)

RE.CNT 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD RE

- (1) Abe, T; J Microelectromech Syst 1998, V7, P94
- (2) Bressers, P; J Electrochem Soc 1996, V143, P1744 CA
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- (4) Gutsche, C; J Am Chem Soc 1981, V103, P3782 CA
- (5) Moldovan, C; Abstract Book workshop of physical chemistry of wet chemical etching of silicon 1998, P21
- (6) Pauling, L; General Chemistry 1970, P461
- (7) Ristic, L; Sensor Technology and Devices P67
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- (11) Shinkai, S; J Chem Soc Perkin Trans 1 1990, P3333 CA
- (12) Tan, S; J Microelectromech Syst 1996, V5, P65
- L3 ANSWER 14 OF 30 CA COPYRIGHT 2002 ACS
- AN 131:66347 CA
- TI Silicon hillocks elimination using a complexant redox alkaline system
- AU Moldovan, Carmen; Iosub, Rodica; Dascalu, Dan; Nechifor, Gheorghe; Danila, Carmen
- CS National Institute for R and D in Microtechnologies, Bucharest, R72225, Rom.
- SO Proceedings of SPIE-The International Society for Optical Engineering (1999), 3680(Pt. 2, Design, Test and Microfabrication of MEMS and MOEMS), 1056-1067

 CODEN: PSISDG; ISSN: 0277-786X
 - SPIE-The International Society for Optical Engineering
- DT Journal

PB

- LA English
- CC 76-3 (Electric Phenomena)
- AB This paper presents the results from the investigation of the chem. anisotropic etching of single-crystal silicon
 .ltbbrac.100.rtbbrac. in the following solns.: KOH (or NaOH,
 LiOH.cntdot.H2O), K3[Fe(CN)6] 0.1M, K4[Fe(CN)6] .cntdot.3H2O 0.1M, KNO3
 0.1M and/or complexant added. The complexants added in KOH soln. were:
 calix[4]arenes, phenols, and ether dibenzo 18 crown 6. Results
 using also NaOH or LiOH.cntdot.H2O and complexants are presented. The
 reaction mechanism and the hillocks formation and elimination are
 analyzed. The results allow us to use the redox system and/or the org.
 complexants, to monitor the etching process, to obtain a smooth
 silicon surface, almost free of hillocks, to utilize the usual mask
 material resistant to the new etchants.
- ST silicon hillock elimination; complexant redox alk system silicon hillock
- IT 7440-21-3, Silicon, processes
 - RL: PEP (Physical, engineering or chemical process); PROC (Process) (silicon hillock elimination using complexant redox alk. system)
- RE.CNT 10 THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD RE
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(6) Ristic, L; Sensor Technology and Devices 1994, P67
(7) Sato, K; Sensors and Actuators A 1998, V64, P87
(8) Seidel, H; Journal of the Electrochemical Society 1990, V137, P3612 CA
(9) Shinkai, S; J Chem Soc Perkin Trans 1 1990, P3333 CA
(10) Tan, S; Journal of Microelectromechanical Systems 1996, V5, P65
    ANSWER 15 OF 30 CA COPYRIGHT 2002 ACS
L3
AN
     130:317109 CA
     Anisotropic etching of silicon in a complexant redox alkaline
TI
     system
    Moldovan, Carmen; Iosub, Rodica; Dascalu, Dan; Nechifor, Gheorghe.
ΑU
     Institute of Microtechnology, Bucharest, R72225, Rom.
CS
     Eurosensors XII, Proceedings of the 12th European Conference on
SO
     Solid-State Transducers and the 9th UK Conference on Sensors and Their
     Applications, Southampton, UK, Sept. 13-16, 1998 (1998), Volume 2,
     1009-1012. Editor(s): White, N. M. Publisher: Institute of Physics
     Publishing, Bristol, UK.
     CODEN: 67PNAZ
DT
     Conference
LΑ
     English
     67-3 (Catalysis, Reaction Kinetics, and Inorganic Reaction Mechanisms)
CC
     This paper presents the results from an investigation of the chem.
AΒ
     anisotropic etching of silicon in the following soln.: KOH 4.5M,
     K3[Fe(CN)6] 0.1M, K4[Fe(CN)6] 3H2O 0.1M, KNO3 0.1M and complexant added.
     The reaction mechanism, the etch rate and the roughness are
     analyzed.
     hydroxide potassium anisotropic etching kinetics silicon;
ST
     ferricyanide potassium anisotropic etching kinetics silicon;
     ferrocyanide potassium anisotropic etching kinetics silicon;
     nitrate potassium anisotropic etching kinetics silicon;
     complexant redox alk anisotropic etching kinetics silicon
IT
     Etching kinetics
        (anisotropic etching of silicon in complexant redox alk.
        system)
ΙT
     Crown ethers
     Metacyclophanes
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); RCT
     (Reactant); PROC (Process); RACT (Reactant or reagent)
        (anisotropic etching of silicon in complexant redox alk.
        system)
IT
     Etching
        (mechanism; anisotropic etching of silicon in complexant
        redox alk. system)
                               1310-58-3, Potassium hydroxide,
     281-54-9, Calix-4-arene
IT
                 7440-21-3, Silicon, reactions 7757-79-1, Potassium nitrate,
     reactions
                 13746-66-2, Potassium ferricyanide 13943-58-3, Potassium
     reactions
     ferrocyanide
                   14187-32-7, Dibenzo-18-crown-6
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); RCT
     (Reactant); PROC (Process); RACT (Reactant or reagent)
        (anisotropic etching of silicon in complexant redox alk.
        system)
              THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
RE
(1) Pauling, L; General Chemistry 1970
(2) Ristic, L; Sensor Technology and Devices 1994, P67
(3) Seidel, H; J Electrochem Soc 1990, V137(11), P3612 CA
    ANSWER 16 OF 30 CA COPYRIGHT 2002 ACS
L3
ΑN
     130:260101 CA
     Mechanism of anisotropic etching of silicon in a complexant
ΤI
     alkaline system
ΑU
     Moldovan, Carmen; Iosub, Rodica; Nechifor, Ghe.; Dascalu, D.; Craciunoiu,
     F.; Serban, B.
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National Institute for Research and Development in Microtechnologies,

CS

Bucharest, R72225, Rom.

- SO CAS '98 Proceedings, International Semiconductor Conference, 21st, Sinaia, Rom., Oct. 6-10, 1998 (1998), Volume 2, 353-356 Publisher: Institute of Electrical and Electronics Engineers, New York, N. Y. CODEN: 67HUAG
- DT Conference
- LA English
- CC 76-3 (Electric Phenomena)
- AB This paper presents the results of the study and expts. of the chem. anisotropic etching of silicon in a complexant alk. system (KOH 4.5M and complexants added). The great results obtained using calix[4] arene like complexant make necessary the study of the mechanism of the silicon etch rate increasing and of the roughness minimizing. A new macrocyclic complexant (azocalix[4] arene) and indicator of pH for silicon etching soln., the etch rate and the roughness are analyzed. The complexant alk. system for anisotropic etching of silicon is an absolutely original idea of the authors.
- ST silicon anisotropic etching cleaning mechanism; alkali system etching silicon semiconductor
- IT Etching

Semiconductor materials

(mechanism of anisotropic **etching** of silicon in a complexant alk. system)

IT 7440-21-3, Silicon, processes

RL: PEP (Physical, engineering or chemical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses)

(mechanism of anisotropic **etching** of silicon in a complexant alk. system)

IT 1310-58-3, Potassium hydroxide (KOH), reactions

RL: RCT (Reactant); RACT (Reactant or reagent)
 (mechanism of anisotropic etching of silicon in a complexant
 alk. system)

- RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD RE
- (1) Luca, C; Aplicatii ale Chimiei Supramoleculare 1996
- (2) Moldovan, C; Anisotropic Etching of Silicon in a Complexant Redox Alkaline System 1998, EUROSENSORS XII
- (3) Moldovan, C; Workshop on Physical Chemistry of Wet Chemical Etching of Silicon 1998
- (4) Ristic, L; Sensor Technology and Devices 1994, P67
- (5) Seidel, H; J Electrochem Soc 1990, V137(11), P3612 CA
- L3 ANSWER 17 OF 30 CA COPYRIGHT 2002 ACS
- AN 130:202831 CA
- TI Sub-10 nm electron beam lithography using a poly(.alpha.-methylstyrene) resist with a molecular weight of 650
- AU Manako, Shoko; Fujita, Jun-Ichi; Tanigaki, Katsumi; Ochiai, Yukinori; Nomura, Eiichi
- CS Fundamental Research Laboratories, NEC Corporation, Tsukuba, 305-8501, Japan
- SO Japanese Journal of Applied Physics, Part 1: Regular Papers, Short Notes & Review Papers (1998), 37(12B), 6785-6787 CODEN: JAPNDE; ISSN: 0021-4922
- PB Japanese Journal of Applied Physics
- DT Journal
- LA English
- CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
- AB Only the 7 nm patterning has been achieved by using an org. neg. electron beam (EB) resist, poly(.alpha.-methylstyrene) resist with a mol. wt. of 650 (.alpha.MST650). An .alpha.MST650 resist film can be prepd. by a conventional spin-coating technique and its surface roughness is smooth, with peak-to-valley roughness of less than 1.5 nm. Although the

sensitivity of .alpha.MST650 is very low, 30 mC/cm2, high-resoln. patterns of 15 nm width and 25 nm pitch with no scum have been fabricated using a 50 kV electron beam of about 7 nm diam. The etching selectivity between Si and .alpha.MST650 is about 2.3 and the etching durability of .alpha.MST650 is superior to those of calixarene resist and polystyrene resist. The min. line pattern sizes achievable with .alpha.MST650 resist is 7 nm, although the min. line pattern size of polystyrene with a mol. wt. of 1100 is about 12 nm. The pattern size dependence of resist resin mol. wt. exit in beyond resist mol. wt. of 650. electron beam lithog polymethylstyrene resist Electron beam resists (sub-10 nm electron beam lithog. using poly(.alpha.-methylstyrene) resist with mol. wt. of 650) 75-73-0, Carbon tetrafluoride RL: NUU (Other use, unclassified); USES (Uses) (plasma etch; sub-10 nm electron beam lithog. using poly(.alpha.-methylstyrene) resist with mol. wt. of 650) 25014-31-7, Poly(.alpha.-methylstyrene) RL: TEM (Technical or engineered material use); USES (Uses) (sub-10 nm electron beam lithog. using poly(.alpha.-methylstyrene) resist with mol. wt. of 650) THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT 9 (1) Beaumont, S; Microelectron Eng 1996, V32, P283 CA (2) Brandrup, J; Polymer Handbook 3rd ed 1989 (3) Chang, T; IBM J Res Develop 1988, V32, P462 CA (4) Chou, S; J Vac Sci Technol B 1997, V15, P2897 CA (5) Fujita, J; J Vac Sci Technol B 1996, V14, P4272 CA (6) Kawaura, H; Ext Abstr 1997 Int Conf Solid State Devices and Materials 1997, P572 (7) Manako, S; Jpn J Appl Phys 1997, V36, P7773 CA (8) Manako, S; Jpn J Appl Phys 1997, V36, PL724 CA (9) Yoshimura, T; Appl Phys Lett 1996, V68, P1799 CA ANSWER 18 OF 30 CA COPYRIGHT 2002 ACS 130:132297 CA 25-nm-pitch (Ga, In) As/InP buried structure: improvement by calixarene as an electron beam resist and tertiarybutylphosphine as a P source in organometallic vapor phase epitaxy regrowth Miyamoto, Y.; Kokubo, A.; Hattori, T.; Hongo, H.; Suhara, M.; Furuya, K. Department of Electrical and Electronic Engineering, Tokyo Institute of Technology, O-okayama, Meguro-ku, Tokyo, 152, Japan Journal of Vacuum Science & Technology, B: Microelectronics and Nanometer Structures (1998), 16(6), 3894-3898 CODEN: JVTBD9; ISSN: 0734-211X American Institute of Physics Journal English 76-3 (Electric Phenomena) To achieve a fine periodic semiconductor structure by electron-beam lithog.; calixarene was used as an electron-beam resist. A 25-nm-pitch InP pattern was formed successfully, and 40-nm-pitch InP structures were achieved with good reproducibility. A shorter developing time, precise stage motion, accurate control of the widths of lines and spaces, and slight O2 ashing were important to obtain a fine InP pattern by a two-step wet chem. etching process. Further, the fabricated periodic InP pattern was buried in a (Ga, In) As structure by organometallic vapor phase epitaxy. The introduction of tertiarybutylphosphine as the phosphorus source prevented the fine structure from deforming when the temp. was raised and a 25-nm-pitch periodic structure was buried successfully. indium phosphide fine periodic structure fabrication Electron beam resists

(calixarene; fabrication of fine periodic InP structure by

ŚТ

IT

IT

IT

RE

L3

AN

TI

ΑU

CS

so

PΒ

DT

LA

CC

AΒ

ST

ΙT

```
electron-beam lithog using calixarene electron-beam resist)
IT
     Electron beam lithography
        (fabrication of fine periodic InP structure by electron-beam lithog
        using calixarene electron-beam resist)
     141137-71-5
IT
     RL: NUU (Other use, unclassified); USES (Uses)
        (electron-beam resist; fabrication of fine periodic InP structure by
        electron-beam lithog using calixarene electron-beam resist)
IT
     22398-80-7, Indium monophosphide, uses
     RL: DEV (Device component use); USES (Uses)
        (fabrication of fine periodic InP structure by electron-beam lithog
        using calixarene electron-beam resist)
              THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 9
RE
(1) Coldren, L; Electron Lett 1982, V18, P235 CA
(2) Fujita, J; Annual Meeting of Japan Society of Applied Physics 1996
(3) Fujita, J; Appl Phys Lett 1996, V68, P1297 CA
(4) Furuya, K; J Appl Phys 1987, V62, P1492
(5) Hongo, H; Jpn J Appl Phys Part 1 1994, V33, P925 CA
(6) Hongo, H; Jpn J Appl Phys Part 1 1998, V37, P1518 CA
(7) Inamura, E; Jpn J Appl Phys Part 1 1989, V28, P2193 CA
(8) Komeno, J; J Cryst Growth 1994, V145, P468 CA
(9) Suhara, M; J Cryst Growth 1994, V145, P698 CA
     ANSWER 19 OF 30 CA COPYRIGHT 2002 ACS
L3
AN
     129:237591 CA
ΤI
     Calixarene resists for nano-lithography
AIJ
     Ohnishi, Yoshitake; Wamme, Naoko; Fujita, Jun-Ichi
CS
     Fundamental Research Laboratories, NEC Corporation, Tsukuba, 305, Japan
SO
     ACS Symp. Ser. (1998), 706 (Micro- and Nanopatterning Polymers), 249-261
     CODEN: ACSMC8; ISSN: 0097-6156
PR
     American Chemical Society
DT
     Journal
LΑ
     English
CC
     74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other
     Reprographic Processes)
AB
     Calixarenes were developed as neg. electron resists for
     nano-lithog. These cluster-like, or roughly ball-shaped mols. form very
     flat and hard films by spin-coating. The high resoln. of these resists
     down to several nm is because these mols. are quite small and free as is
     in ordinary chain polymers. As etching resistance of
     calixarenes is sufficient in plasma-etch processes,
     nano-fabrication of metal or semiconductors is easily carried out by
     conventional resist processes.
ST
     calixarene resist nanolithog
IT
     Clusters
       Etching
     Lithography
     Mass spectra
     Resists
     Spin coating
        (calixarene resists for nanolithog.)
IT
     Metacyclophanes
     RL: DEV (Device component use); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (calixarene resists for nanolithog.)
IT
     124006-38-8
                   141137-71-5
     RL: DEV (Device component use); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (calixarene resists for nanolithog.)
L3
     ANSWER 20 OF 30 CA COPYRIGHT 2002 ACS
AN
     129:209993 CA
ΤI
     A 25-nm-pitch GaInAs/InP buried structure using calixarene
```

resist ΑU Kokubo, Atsushi; Hattori, Etsuya; Hongo, Hiroo; Suhara, Michihiko; Miyamoto, Yasuyuki; Furuya, Kazuhito CS Department of Electrical and Electronic Engineering, Tokyo Institute of Technology, Okayanal, Meguro-ku, Tokyo, 152, Japan SO Jpn. J. Appl. Phys., Part 2 (1998), 37(7A), L827-L829 CODEN: JAPLD8; ISSN: 0021-4922 PB Japanese Journal of Applied Physics Journal דת English LА CC 76-3 (Electric Phenomena) Section cross-reference(s): 74 ΔR To realize a fine periodical pattern by electron beam lithog., a study for using calixarene as a resist was carried out. A 25-nm-pitch resist pattern was fabricated and transferred to a thin InP layer by 2-step wet chem. etching. Precise slight 02 ashing, to eliminate residual matter was essential to transfer the pattern by wet etching. The controllability of the width was improved when using calixarene, when the period was 40 nm. Also, a 25-nm-pitch InP pattern was buried in a GaInAs structure by OMVPE. This technol. could be applied to realize electron wave devices. ST gallium indium arsenide phosphide calixarene resist; buried structure calixarene resist arsenide phosphide TT Electron beam lithography Etching Metalorganic vapor phase epitaxy Plasma ashing Resists (a 25-nm-pitch GaInAs/InP buried structure using calixarene resist) IT Metacyclophanes RL: DEV (Device component use); USES (Uses) (a 25-nm-pitch GaInAs/InP buried structure using calixarene resist) IT 22398-80-7, Indium phosphide, properties 106070-25-1, Gallium indium arsenide RL: DEV (Device component use); PRP (Properties); USES (Uses) (a 25-nm-pitch GaInAs/InP buried structure using calixarene resist) ANSWER 21 OF 30 CA COPYRIGHT 2002 ACS L3 128:187114 CA AN Single-electron transistors fabricated from a doped-Si film in a TIsilicon-on-insulator substrate ΑU Sakamoto, T.; Kawaura, H.; Baba, T. CS NEC Fundamental Research Laboratories, Tsukuba, Ibaraki, 305, Japan SO Appl. Phys. Lett. (1998), 72(7), 795-796 CODEN: APPLAB; ISSN: 0003-6951 PB American Institute of Physics DТ Journal LA English CC 76-3 (Electric Phenomena) AB The authors propose doped-thin-Si-film single-electron transistors (DS-SETs), which are fabricated from a highly doped Si film in a Si-on-insulator substrate by electron-beam lithog. with a high-resoln. resist (calixarene) and dry etching with CF4 gas. Because the structure can be well controlled, the DS-SET with a 45-nm-diam. island shows nearly ideal characteristics of SETs with a charging energy of 1.4 meV. Single-electron tunneling occurs through a single island without any isolated islands formed in potential

fluctuations. The authors also discuss the discreteness of energy levels

ST single electron transistor doped silicon film

IT Dry etching

in a Si island.

```
Electron beam lithography
        (single-electron transistors fabricated from doped-Si film in
        silicon-on-insulator substrate by)
IT
     Transistors
        (single-electron; single-electron transistors fabricated from doped-Si
        film in silicon-on-insulator substrate)
IT
     7440-21-3, Silicon, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (single-electron transistors fabricated from doped-Si film in
        silicon-on-insulator substrate)
TΤ
     75-73-0, Carbon fluoride (CF4)
     RL: TEM (Technical or engineered material use); USES (Uses)
        (single-electron transistors fabricated from doped-Si film in
        silicon-on-insulator substrate by dry etching with)
     ANSWER 22 OF 30 CA COPYRIGHT 2002 ACS
L3
     128:147408 CA
AN
     Calixarene electron beam resist for nano-lithography
TI
     Fujita, Jun-ichi; Ohnishi, Yoshitake; Manako, Shoko; Ochiai, Yukinori;
ΑU
     Nomura, Eiichi; Sakamoto, Toshitsugu; Matsui, Shiniji
     Fundamental Res. Lab., NEC Corp., Tsukuba, 305, Japan
CS
     Jpn. J. Appl. Phys., Part 1 (1997), 36(12B), 7769-7772
SO
     CODEN: JAPNDE; ISSN: 0021-4922
PB
     Japanese Journal of Applied Physics
DT
     Journal
LA
     English
     74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other
CC
     Reprographic Processes)
     New electron beam (EB) resists made of calixarene resists are
AB
     introduced. Typical sensitivities of calixarene resists range
     from 700 .mu./C/cm2 to 7 mC/cm2. High-d. dot arrays with 15 nm diam.
     constructed using calixarene resist were easily delineated using
     a point EB lithog. system. Our results suggest that the resoln. limit of
     calixarene resists is dominated by a development process such as
     adhesion to a substrate rather than by the EB profile. Calixarene
     resists are resistant to etching by halide plasma. We also
     demonstrated nanoscale devices processed by using calixarene
     resists. Calixarene resists are promising materials for
     nanofabrication.
ST
     calixarene electron beam resist nano lithog
IT
     Electron beam resists
        (calixarene; for nano-lithog.)
TΤ
     Metacyclophanes
     RL: PRP (Properties); TEM (Technical or engineered material use); USES
     (Uses)
        (electron beam resist for nano-lithog.)
IT
     Electron beam lithography
        (nano-; calixarene electron beam resist for)
TT
     124006-38-8
                   141137-71-5
     RL: PRP (Properties); TEM (Technical or engineered material use); USES
     (Uses)
        (electron beam resist for nano-lithog.)
     ANSWER 23 OF 30 CA COPYRIGHT 2002 ACS
L3
AN
     128:121581 CA
ΤI
     Process optimization for production of sub-20 nm soft x-ray zone plates
     Spector, S. J.; Jacobsen, C. J.; Tennant, D. M.
ΑU
     Department of Physics, SUNY at Stony Brook, Stony Brook, NY, 11794, USA
CS
SO
     J. Vac. Sci. Technol., B (1997), 15(6), 2872-2876
     CODEN: JVTBD9; ISSN: 0734-211X
PB
     American Institute of Physics
DT
     Journal
T.A
     English
CC
     74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other
```

Reprographic Processes)
Section cross-reference(s): 73

- AΒ We report here the optimization of processes for producing sub-20 nm soft x-ray zone plates, using a general purpose electron beam lithog. system and com. resist technologies. We have critically evaluated the failure point of the various process steps and where possible chosen alternate methods, materials, or otherwise modified the process. Advances have been made in most steps of the process, including the imaging resist, pattern conversion for electron beam exposure, and pattern transfer. Two phase shifting absorber materials, germanium and nickel, were compared. Zone plates with 30 nm outer zones have been fabricated in both germanium and nickel with excellent quality using PMMA and zones as small as 20 nm have been fabricated in nickel using the calixarene resist. The total efficiency as well as the efficiency of different regions of the zone plates were measured. All zone plates have demonstrated good efficiencies, with nickel zone plates performing better than germanium zone plates.
- ST x ray zone plate lithog fabrication
- IT X-ray devices

(lenses; process optimization for electron lithog. prodn. of sub-20 nm soft x-ray zone plates)

IT Electron beam lithography

Electron beam resists

(process optimization for electron lithog. prodn. of $sub-20\ nm\ soft\ x-ray\ zone\ plates)$

IT Lenses

(x-ray; process optimization for electron lithog. prodn. of sub-20 nm soft x-ray zone plates)

IT Optical instruments

(zone plates, Fresnel, x-ray; process optimization for electron lithog. prodn. of sub-20 nm soft x-ray zone plates)

TT 75-63-8, Trifluorobromomethane 7782-44-7, Oxygen, uses
RL: NUU (Other use, unclassified); USES (Uses)
 (etchant; process optimization for electron lithog. prodn. of
 sub-20 nm soft x-ray zone plates)

IT 12033-89-5; Silicon nitride, processes

RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)

(process optimization for electron lithog. prodn. of sub-20 nm soft x-ray zone plates)

IT 9011-14-7, PMMA 89072-52-6, AZ 4110

RL: PEP (Physical, engineering or chemical process); PROC (Process) (process optimization for electron lithog. prodn. of sub-20 nm soft x-ray zone plates)

(shifting absorber; process optimization for electron lithog. prodn. of sub-20 nm soft x-ray zone plates)

- L3 ANSWER 24 OF 30 CA COPYRIGHT 2002 ACS
- AN 127:314181 CA
- TI Strategies toward the development of integrated chemical sensors fabricated from light emitting porous silicon
- AU Coffer, Jeffery L.; Zhang, Libing; John, John St.
- CS Department of Chemistry, Texas Christian University, Ft. Worth, TX, 76129, USA
- SO Proc. SPIE-Int. Soc. Opt. Eng. (1997), 3226 (Microelectronic Structures and MEMS for Optical Processing III), 168-179 CODEN: PSISDG; ISSN: 0277-786X

```
PB
     SPIE-The International Society for Optical Engineering
DT
     Journal
     English
LA
     79-2 (Inorganic Analytical Chemistry)
CC
     Section cross-reference(s): 73, 76, 80
AB
     Several different approaches designed to produce an opto-electronic chem.
     sensor based on light-emitting porous Si are described, all of which
     entail modification of the as-formed porous Si surface to alter device
     characteristics. The issue of selectivity and sensitivity of a given
     porous Si sensor can be modified by coating the porous Si surface with
     basket-shaped mols. known as calixarenes; the ability of such a
     structure to detect copper ions and organonitrogen compds. is reported.
     Surface modification of porous Si through etching and deposition
     of conducting polymers to alter Si light emission color and intensity is
     also discussed. The fabrication of porous Si-based waveguides on Si and
     the impact of surface modification with erbium ions are also described.
     integrated sensor light emitting porous silicon; copper detection light
ST
     emitting silicon sensor; amine detection light emitting silicon sensor;
     calixarene coating light emitting silicon sensor; conducting
     polymer coating light emitting silicon; erbium effect porous silicon
     wavequide
     Waveguides
IT
        (fabrication of porous Si-based waveguides on porous Si and impact of
        surface modification with erbium ions)
     Electroluminescent devices
ΙT
        (integrated chem. sensors fabricated from light emitting porous
        silicon)
ΙT
     Amines, analysis
     RL: ANT (Analyte); PRP (Properties); ANST (Analytical study)
        (integrated chem. sensors fabricated from light emitting porous silicon
        for detection of)
IT
     Luminescence
        (photoluminescence quenching of calixarene coated porous
        silicon by Cu(II) and amines)
ΙT
     Optical sensors
        (semiconductive luminescence; integrated chem. sensors fabricated from
        light emitting porous silicon)
IT
     Electroluminescence
        (tunability; of conducting polymer coated porous silicon for chem.
        sensors)
IT
     11106-95-9
     RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (contact for porous Si light emitting diodes)
IT
     7440-52-0, Erbium, uses
     RL: NUU (Other use, unclassified); PRP (Properties); USES (Uses)
        (fabrication of porous Si-based wavequides on porous Si and impact of
        surface modification with erbium ions)
IT
     7440-21-3, Silicon, analysis
     RL: ARU (Analytical role, unclassified); DEV (Device component use); ANST
     (Analytical study); USES (Uses)
        (integrated chem. sensors fabricated from light emitting porous
        silicon)
IT
     25067-59-8, Poly(9-vinyl carbazole)
                                           190785-27-4
     RL: ARU (Analytical role, unclassified); DEV (Device component use); PRP
     (Properties); ANST (Analytical study); USES (Uses)
        (integrated chem. sensors fabricated from light emitting porous silicon
        coated with)
ΙT
     107-10-8, 1-Propanamine, analysis
                                        109-73-9, 1-Butanamine, analysis
     110-58-7, n-Pentylamine
                              7440-50-8, Copper, analysis
     RL: ANT (Analyte); PRP (Properties); ANST (Analytical study)
        (integrated chem. sensors fabricated from light emitting porous silicon
        for detection of)
                   119707-72-1 119707-73-2 127594-37-0
IT
     119707-70-9
                                                             185628-43-7
```

197298-64-9

RL: ARU (Analytical role, unclassified); DEV (Device component use); ANST (Analytical study); USES (Uses)

(integrated chem. sensors fabricated from light emitting porous silicon modified with coating contg.)

- L3 ANSWER 25 OF 30 CA COPYRIGHT 2002 ACS
- AN 127:285942 CA
- TI Super-fine pattern formation and super-fine **etching** method using resist containing **calixarene**
- IN Onishi, Yoshitake; Fujita, Junichi; Aldoeni, Arturo; Casnati, Alessandro; Pochini, Andrea; Ungaro, Rocco
- PA NEC Corp., Japan
- SO Jpn. Kokai Tokkyo Koho, 8 pp.

CODEN: JKXXAF

- DT Patent
- LA Japanese
- IC ICM G03F007-038
 - ICS C09D165-00; C23F001-00; H01L021-027; H01L021-302
- CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

FAN.CNT 1

TAN CONT					
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
1	PI JP 09236919	A2	19970909	JP 1996-157287	19960618
	JP 2792508	B2	19980903		
	US 5702620	Α	19971230	US 1996-693672	19960813
1	PRAT TT 1996-MT382		19960228		

- The title patterning method comprises the steps of forming a resist film made of 5,11,17,23,29,35-hexachloromethyl-37,38,39,40,41,42-hexamethoxycalix[6] arene (I) which is sol. in solvents and sensitive toward high energy rays, exposing the film selectively with the rays, and removing the unexposed area with the solvents to develop the exposed area. The title etching method comprises forming a pattern on a substrate by the above process and dry-etching the substrate along with the pattern. Super-fine resist patterns of the order of nano-meters are obtained. Thus, a soln. of I in dichlorobenzene was coated on a Si wafer, pre-baked, patternwise exposed with an electron beam, and developed with xylene to form a pattern.
- ST calixarene neg working resist pattern formation; etching calixarene neg working resist
- IT Resists

(etching; super-fine pattern formation using resist contg.
calixarene)

IT Metacyclophanes

RL: PNU (Preparation, unclassified); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(super-fine pattern formation using resist contg. calixarene)

IT 106750-73-6P

RL: PNU (Preparation, unclassified); RCT (Reactant); PREP (Preparation) (prepn. of chloromethylmethoxcalixarene)

IT 74-88-4, Methyl iodide, reactions 24566-90-3, Chloromethyl octyl ether 96627-08-6, Calix[6] arene

RL: RCT (Reactant)

(prepn. of chloromethylmethoxcalixarene)

IT 124006-38-8P

RL: PNU (Preparation, unclassified); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)
(super-fine pattern formation using resist contg. calixarene)

- L3 ANSWER 26 OF 30 CA COPYRIGHT 2002 ACS
- AN 126:150389 CA
- TI Nanometer-scale resolution of **calixarene** negative resist in electron beam lithography

```
AU Fujita, J.; Ohnishi, Y.; Ochiai, Y.; Nomura, E.; Matsui, S.
```

- CS Fundamental Research Laboratories, NEC Corporation, Tsukuba, 305, Japan
- SO J. Vac. Sci. Technol., B (1996), 14(6), 4272-4276 CODEN: JVTBD9; ISSN: 0734-211X
- PB American Institute of Physics
- DT Journal
- LA English
- CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
- AB New non-polymer materials, calixarene derivs. were tested as high-resoln. neg. resists for use in electron beam lithog. Arrays of 12-nm-diam. dots with a 25 nm pitch were fabricated easily. The sensitivity of calixarene in terms of area dose ranged from 700 to 7000 .mu.C/cm2, and the required dose for dot fabrication was about 105 electrons/dot. The std. area dose for calixarene is almost 20 times higher than that for polymethyl methacrylate (PMMA), but the electron spot dose for dot fabrication by calixarene is almost the same as that for PMMA and other highly sensitive resists such as SAL (chem. amplified neg. resist for electron beam made by Shipley). The electron spot dose for such extremely small dots does not seem to depend on std. area dose, but any resist tends to require the same dose under exposure in a 50 keV electron beam writing system. We propose a qual. exposure model that suggests a tradeoff of dose and dot size. calixarene seems to be promising material for nanofabrication.
- ST calixarene neg resist electron beam lithog
- IT Resists

(neg.-working; non-polymer calixarene deriv. as high-resoln.
neg. resist for electron beam lithog.)

IT Electron beam lithography

Etching

(non-polymer calixarene deriv. as high-resoln. neg. resist for electron beam lithog.)

IT Metacyclophanes

RL: PEP (Physical, engineering or chemical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses)

(non-polymer calixarene deriv. as high-resoln. neg. resist for electron beam lithog.)

IT 124006-38-8 141137-71-5, 5,11,17,23,29,35-Hexamethyl-37,38,39,40,41,42hexaacetoxycalix[6] arene

RL: PEP (Physical, engineering or chemical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses)

(non-polymer calixarene deriv. as high-resoln. neg. resist for electron beam lithog.)

- L3 ANSWER 27 OF 30 CA COPYRIGHT 2002 ACS
- AN 124:215824 CA
- TI Ultrahigh resolution of **calixarene** negative resist in electron beam lithography
- AU Fujita, J.; Ohnishi, Y.; Ochiai, Y.; Matsui, S.
- CS Fundam. Res. Lab., NEC Corp., Tsukuba, 305, Japan
- SO Appl. Phys. Lett. (1996), 68(9), 1297-9 CODEN: APPLAB; ISSN: 0003-6951
- DT Journal
- LA English
- CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
- AB A nonpolymer material, calixarene deriv. (hexaacetate p-methylcalix[6]arene) was tested as a high-resoln. neg. resist under an electron beam lithog. process. It showed under 10-mm resoln. with little side roughness and high durability to halide plasma etching. A sub-10-nm Ge quantum wire was perfectly etched off without defects. Such a performance is suitable for nanoscale device processes.
- ST electron beam lithog calixarene neg resist; hexaacetate methylcalixarene neg resist electron lithog

```
(FILE 'HOME' ENTERED AT 10:07:40 ON 10 MAY 2002)
    FILE 'CA' ENTERED AT 10:07:45 ON 10 MAY 2002
L1
             6 S DENTRIT? AND ETCH?
L2
             30 S CALIX? AND ETCH?
L3
            30 S L2 NOT L1
L4
            236 S CALIX? AND PHOTO?
L5
             50 S L4 AND HYDROX?
L6
             50 S L5 NOT L3
=> s 16 and glycoluril
           357 GLYCOLURIL
L7
             0 L6 AND GLYCOLURIL
=> s 16 and glycouril
            33 GLYCOURIL
             0 L6 AND GLYCOURIL
L8
=> s 16 and powderlink
            55 POWDERLINK
L9
             0 L6 AND POWDERLINK
=> d his
     (FILE 'HOME' ENTERED AT 10:07:40 ON 10 MAY 2002)
     FILE 'CA' ENTERED AT 10:07:45 ON 10 MAY 2002
L1
             6 S DENTRIT? AND ETCH?
             30 S CALIX? AND ETCH?
L2
            30 S L2 NOT L1
L3
           236 S CALIX? AND PHOTO?
L4
             50 S L4 AND HYDROX?
L5
L6
             50 S L5 NOT L3
             0 S L6 AND GLYCOLURIL
L7
             0 S L6 AND GLYCOURIL
L8
             0 S L6 AND POWDERLINK
L9
=> log y
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STN INTERNATIONAL LOGOFF AT 10:10:51 ON 10 MAY 2002